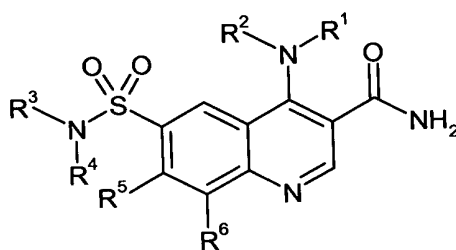


Amendments To The Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

What is claimed is:

1. (Currently Amended) A compound of formula (I) or a pharmaceutically acceptable salt thereof:



(I)

wherein:

R¹ is

Aryl optionally substituted by one or more substituents selected from C₁₋₆alkyl, C₁₋₆alkoxy, halogen, C₁₋₆alkylCO-, -(CH₂)_mOH, -CN, R⁷R⁸N-;

Aryl fused to a C₄₋₇cycloalkyl ring;

Aryl fused to a heterocyclyl ring;

Heteroaryl wherein the heteroaryl is optionally substituted by one or more substituents selected from: C₁₋₆alkyl, N-oxide, C₁₋₆alkoxy; or

Heterocyclyl.

R² is hydrogen or C₁₋₆alkyl;

R³ is

Hydrogen;

C₁₋₆alkyl optionally substituted by one or more substituents selected from: heterocyclyl (itself optionally substituted by C₁₋₆alkyl), R⁹R¹⁰NCO-, R¹¹CONR¹²-, C₁₋₆alkylSO₂NR¹³-, C₁₋₆alkoxy, R¹⁴R¹⁵N-;

C₃₋₇cycloalkyl;

Aryl or aryl(C₁₋₆alkyl) wherein the aryl is optionally substituted by one or more substituents selected from: C₁₋₆alkyl, C₁₋₆alkoxy, halogen, R¹⁶R¹⁷NCO-;

Aryl fused to C₄₋₇cycloalkyl, wherein the cycloalkyl is optionally substituted by =O;

Heteroaryl or heteroaryl(C₁₋₆alkyl), wherein the heteroaryl is optionally substituted by one or more substituents selected from C₁₋₆alkyl, C₁₋₆alkoxy, halogen; or

Heterocyclyl optionally substituted by one or more C₁₋₆alkyl, C₁₋₆alkylCO-, C₁₋₆alkylSO₂-, R¹⁸R¹⁹NCO-, C₁₋₆alkoxyCO-;

R⁴ is hydrogen or C₁₋₆alkyl;

R³ and R⁴ together with the nitrogen atom to which they are attached may form a heterocyclyl ring, which is optionally substituted by one or more substituents selected from C₁₋₆alkyl (optionally substituted by one or more OH or C₁₋₆alkoxy groups), C₁₋₆alkoxy, C₁₋₆alkoxyCO-, C₃₋₇cycloalkyl (optionally substituted by OH), C₁₋₆alkylCO-, C₁₋₆alkylSO₂-, OH, -(CH₂)_mNR²⁰R²¹, -(CH₂)_mCONR²²R²³, -(CH₂)_mNR²⁴COR²⁵, C₁₋₆alkoxyC₁₋₄alkyl, arylCO- heteroaryl, heteroarylC₁₋₄alkyl, heteroarylCO.

m is 0-6

R⁵ is hydrogen or C₁₋₆alkyl;

R⁶ is hydrogen, C₁₋₆alkyl, C₁₋₆alkoxy, fluorine, chlorine, or bromine;

R⁷⁻²⁵ all independently represent hydrogen, or C₁₋₆ alkyl;

R¹⁴ and R¹⁵ together with the nitrogen atom to which they are attached may form a heterocyclyl ring;

R¹⁶ and R¹⁷ together with the nitrogen atom to which they are attached may form a heterocyclyl ring;

R¹⁸ and R¹⁹ together with the nitrogen atom to which they are attached may form a heterocyclyl ring;

R²⁰ and R²¹ together with the nitrogen atom to which they are attached may form a heterocyclyl ring; and

R²² and R²³ together with the nitrogen atom to which they are attached may form a heterocyclyl ring.

2. (Currently Amended) A compound according to claim 1 wherein R¹ is selected from

aryl optionally substituted by one or more substituents selected from C₁₋₆alkyl, C₁₋₆alkoxy-, halogen, -CN;

aryl fused to a heterocyclyl ring; and

heteroaryl optionally substituted by one or more substituents selected from: C₁₋₆alkyl.

3. (Currently Amended) A compound according to claim 1 ~~or 2~~ wherein R^2 is hydrogen.

4. (Currently Amended) A compound according to ~~any of claims 1 to 3~~ wherein R^3 is selected from

C_{1-6} alkyl optionally substituted by one or more substituents selected from heterocyclyl, C_{1-6} alkoxy;

C_{3-7} cycloalkyl; and

Heterocyclyl.

5. (Currently Amended) A compound according to ~~any of claims 1 to 4~~ wherein R^4 is hydrogen or C_{1-6} alkyl.

6. (Currently Amended) A compound according to ~~any of claims 1 to 3~~ wherein R^3 and R^4 together with the nitrogen atom to which they are attached may form a heterocyclyl ring, optionally substituted by one or more substituents selected from C_{1-6} alkyl (optionally substituted by one or more C_{1-6} alkoxy groups), C_{1-6} alkylCO, C_{1-6} alkylSO₂, $-(CH_2)_mCONR^{22}R^{23}$, $-(CH_2)_mNR^{20}R^{21}$, heteroaryl.

7. (Currently Amended) A compound according to ~~any of claims 1 to 6~~ wherein R^5 is hydrogen.

8. (Currently Amended) A compound according to ~~any of claims 1 to 7~~ wherein R^6 is hydrogen or C_{1-6} alkyl.

9. (Currently Amended) A compound according to claim 1 wherein

R^1 is selected from

phenyl optionally substituted by one or more substituents selected from methyl, methoxy, fluoro, chloro, cyano;
dihydrobenzofuranyl; and
indazolyl or benzimidazolyl optionally substituted by methyl;

R² is hydrogen;

R³ is selected from
C₁₋₃alkyl optionally substituted by one C₁₋₂alkoxy group or a 5 to 7 membered saturated ring containing one or two heteratoms selected from nitrogen or oxygen;
C₃₋₅cycloalkyl; and
5 to 7 membered saturated ring containing one heteroatom which is oxygen;

R⁴ is hydrogen or C₁₋₆alkyl;

R⁵ is hydrogen;

R⁶ is hydrogen or C₁₋₆alkyl.

10. (Currently Amended) A compound according to claim 1 wherein

R¹ is selected from

phenyl optionally substituted by one or more substituents selected from methyl, methoxy, fluoro, chloro, cyano;
dihydrobenzofuranyl; and
indazolyl or benzimidazolyl optionally substituted by methyl;

R² is hydrogen;

R³ and R⁴ together with the nitrogen atom to which they are attached may form a 5 or 6 membered heterocyclyl ring, optionally substituted by one or more substituents selected from C₁₋₃alkyl (optionally substituted by one or more C₁.

alkoxy groups), C₁₋₃alkylCO, C₁₋₃alkylSO₂, -CON(CH₃)₂, -N(CH₃)₂, pyrazinyl, pyridinyl;

R⁵ is hydrogen; and

R⁶ is hydrogen or C₁₋₆alkyl.

11. (Previously Presented) A compound of formula (I) selected from the group consisting of

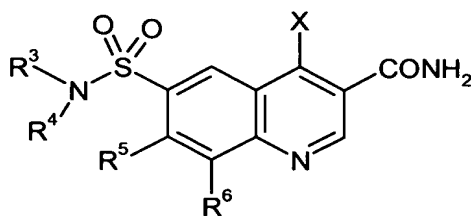
6-[(dimethylamino)sulfonyl]-4-[[3-(methyloxy)phenyl]amino]-3-quinolinecarboxamide;
4-(2,3-dihydro-1-benzofuran-4-ylamino)-6-(4-morpholinylsulfonyl)-3-quinolinecarboxamide;
6-[(4-acetyl-1-piperazinyl)sulfonyl]-4-[[4-fluoro-3-(methyloxy)phenyl]amino]-3-quinolinecarboxamide;
4-[[4-fluoro-3-(methyloxy)phenyl]amino]-6-[[4-(methylsulfonyl)-1-piperazinyl]sulfonyl]-3-quinolinecarboxamide;
6-[(4-acetyl-1-piperazinyl)sulfonyl]-4-(2,3-dihydro-1-benzofuran-4-ylamino)-3-quinolinecarboxamide;
4-(2,3-dihydro-1-benzofuran-4-ylamino)-6-[[4-(methylsulfonyl)-1-piperazinyl]sulfonyl]-3-quinolinecarboxamide;
4-(2,3-dihydro-1-benzofuran-4-ylamino)-6-[(dimethylamino)sulfonyl]-3-quinolinecarboxamide;
6-({4-[(dimethylamino)carbonyl]-1-piperazinyl}sulfonyl)-4-[[4-fluoro-3-(methyloxy)phenyl]amino]-3-quinolinecarboxamide;
4-(2,3-dihydro-1-benzofuran-4-ylamino)-6-[[4-(2-pyrazinyl)-1-piperazinyl]sulfonyl]-3-quinolinecarboxamide;
4-(2,3-dihydro-1-benzofuran-4-ylamino)-6-({4-[(dimethylamino)carbonyl]-1-piperazinyl}sulfonyl)-3-quinolinecarboxamide;
4-(2,3-dihydro-1-benzofuran-4-ylamino)-6-[(tetrahydro-2H-pyran-4-ylamino)sulfonyl]-3-quinolinecarboxamide;
4-[[4-fluoro-3-(methyloxy)phenyl]amino]-8-methyl-6-(4-morpholinylsulfonyl)-3-quinolinecarboxamide

4-(2,3-dihydro-1-benzofuran-4-ylamino)-8-methyl-6-(4-morpholinylsulfonyl)-3-quinolinecarboxamide
8-methyl-4-[(3-methylphenyl)amino]-6-(4-morpholinylsulfonyl)-3-quinolinecarboxamide
4-[(3-fluorophenyl)amino]-8-methyl-6-(4-morpholinylsulfonyl)-3-quinolinecarboxamide
4-[(3-cyanophenyl)amino]-8-methyl-6-(4-morpholinylsulfonyl)-3-quinolinecarboxamide
4-(2,3-dihydro-1-benzofuran-4-ylamino)-6-[[4-(dimethylamino)-1-piperidinyl]sulfonyl]-3-quinolinecarboxamide
4-[(3-chlorophenyl)amino]-8-methyl-6-(4-morpholinylsulfonyl)-3-quinolinecarboxamide
8-methyl-4-[(1-methyl-1H-indazol-6-yl)amino]-6-(4-morpholinylsulfonyl)-3-quinolinecarboxamide
6-[(4-acetyl-1-piperazinyl)sulfonyl]-8-methyl-4-[(3-methylphenyl)amino]-3-quinolinecarboxamide
6-[(4-acetyl-1-piperazinyl)sulfonyl]-4-[[4-fluoro-3-(methyloxy)phenyl]amino]-8-methyl-3-quinolinecarboxamide
6-[(4-acetyl-1-piperazinyl)sulfonyl]-4-(2,3-dihydro-1-benzofuran-4-ylamino)-8-methyl-3-quinolinecarboxamide

and pharmaceutically acceptable salts thereof.

12. (Currently Amended) A process for the preparation of a compound of formula (I) and pharmaceutically acceptable salts thereof as defined in ~~any of~~ claims 1 to 14 which comprises:

(A) reacting a compound of formula (II);



(II)

wherein R³, R⁴, R⁵ and R⁶ are as defined above, and X represents a halogen atom, with an amine of formula R¹R²NH, wherein R¹ and R² are as defined above; or

(B) interconversion of a compound of formula (I) into another compound of formula (I); or

(C) deprotecting a protected derivative of a compound of formula (I).

13.-14. (Canceled).

15. (Currently Amended) A method of treating an inflammatory and/or allergic disease in a mammal in need thereof, which comprises administering to the mammal a therapeutically effective amount of a compound of formula (I) according to ~~any of claims 1 to 14~~, or a pharmaceutically acceptable salt thereof.

16. (Currently Amended) A pharmaceutical composition which comprises a compound according to ~~any of claims 1 to 14~~, or a pharmaceutically acceptable salt thereof ~~optionally~~ with a pharmaceutically acceptable carrier or excipient.

17. (Previously Presented) A pharmaceutical composition according to claim 16 which is suitable for inhaled administration.

18. (Previously Presented) A pharmaceutical composition according to claim 16 which is suitable for oral administration.

Attorney Docket No.: PB60514USw

19. (Previously Presented) A pharmaceutical composition according to claim 16 which is suitable for topical administration.

20. (New) A method of inhibiting PDE4, comprising the administration of the compound of claim 1 or a pharmaceutically acceptable salt thereof.